



(12) **EUROPEAN PATENT APPLICATION**

(43) Date of publication:
16.08.2000 Bulletin 2000/33

(51) Int. Cl.⁷: **B01D 53/94**, F01N 3/10,
B01J 23/40, B01J 23/42

(21) Application number: 00300557.6

(22) Date of filing: 26.01.2000

(84) Designated Contracting States:
**AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU
MC NL PT SE**
Designated Extension States:
AL LT LV MK RO SI

(72) Inventors:
• Jarvis, Mottlene
PA 19352-8933 (US)
• Adams, Karen Marie
Dearborn Heights, Michigan 48125 (US)

(30) Priority: 09.02.1999 US 248189

(71) Applicant:
Ford Global Technologies, Inc.
Dearborn, Michigan 48126 (US)

(74) Representative:
Messulam, Alec Moses et al
A. Messulam & Co. Ltd.,
43-45 High Road
Bushey Heath, Herts WD2 1EE (GB)

(54) **Apparatus and method for treating exhaust gases from a diesel engine using nitrogen oxide absorbent**

(57) The invention is a method and system for treating exhaust gases generated by a diesel engine by locating two catalyst components in the engine exhaust gas passage. The first catalyst component which is exposed to oxidizing diesel exhaust is located nearest to the engine and is a nitrogen oxide absorbent made of support material carrying precious metal. The second component is a catalyst such as a lean-NO_x catalyst or

a selective reduction catalyst which is capable of converting the exhaust gas passing over it including reducing the nitrogen oxides desorbed from the first component into nitrogen (N₂) or nitrous oxide (N₂O). Materials like hydrocarbons or ammonia or urea may be injected into the vicinity of the second catalyst component to aid in the reduction.

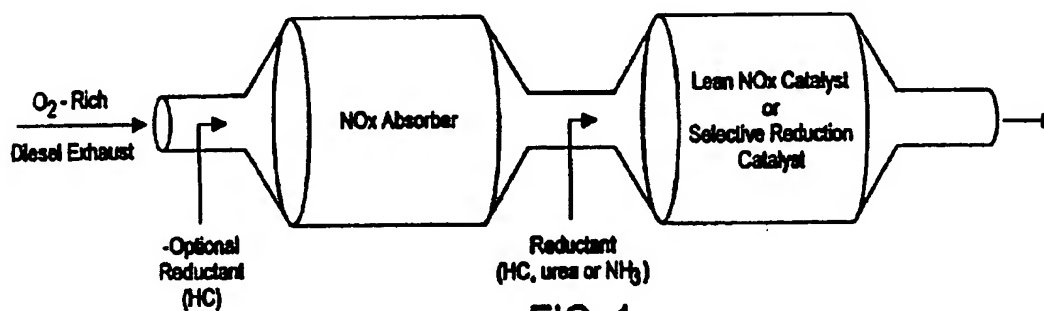


FIG. 1

diesel exhaust gases pass through the first component with NOx being absorbed at low temperatures and later desorbed at elevated temperatures to be reduced over the second component.

[0010] A method and system embodying the present invention is capable of efficiently reducing nitrogen oxides as well as oxidizing hydrocarbons and carbon monoxide in the relatively cool oxidizing conditions of diesel exhaust gases.

[0011] For optimum operation, the amount of nitrogen oxide absorbed on the first component may be monitored so that the NOx absorbed on the first component material can be desorbed by elevating the temperature before the material has reached capacity for nitrogen oxide absorption. Advantageously, however, diesel exhaust temperatures during typical driving, exemplified in the MVEuro cycle, are within the range of NOx desorption often enough that NOx storage capacity in the first component is not likely to be reached.

[0012] The invention will now be described, by way of example, with reference to the accompanying drawings, in which:-

Fig. 1 is a schematic of the invention;

Fig. 2 is a graph showing the NOx storage efficiency and release as a function of gas sample temperature for a Pt/alumina first component embodiment according to the present invention;

Fig. 3 and Fig. 4 are graphs showing NOx storage and conversion efficiency for the first component materials of Fig. 1 using increased hydrocarbon content in the gas sample;

Fig. 5 is a graph showing NOx conversion efficiency over a lean-NOx catalyst useful in an embodiment of the present invention as the second component; and

Fig. 6 is a graph showing NOx storage and conversion of released NOx for a two-component system according to an embodiment of the present invention.

[0013] The invention method comprises locating two catalyst components in the exhaust passage of a diesel engine exhaust gas passage, the first component being located closer to the engine manifold. The first component is a material which, during diesel engine operation, is exposed to exhaust gases which are always oxidizing. During the relatively cool operating temperatures as during engine start up, light acceleration and low speed-load cruises, as particularly takes place during urban driving, the NOx present in the oxidizing exhaust is absorbed on the first component material. Later when the first component is exposed to higher temperatures as during increased load of urban and highway driving, the absorbed NOx desorbs from the first component in the presence of the oxidizing exhaust and is carried in the gas stream to the second component for reduction as over a lean-NOx catalyst or SCR

catalyst. The first component can be positioned to contact engine out gases immediately after they exit e.g., the exhaust manifold, or a turbocharger, or other exhaust after-treatment equipment such as a particulate filter. A schematic of this invention is shown in Fig. 1.

[0014] We have found that, e.g., platinum on alumina stores and releases NOx under O₂ rich conditions, and at exhaust gas temperatures typical of diesel light-duty vehicle driving, e.g., on a standard drive cycle such as MVEuro. This NOx storage occurs over the same low temperatures where Pt is active for lean NOx reduction. Operation of lean-NOx catalysts on diesels typically requires reductant added to the exhaust gases because engine out HC levels are too low. When no HC is present, Pt/alumina stores NOx. When HC is added, both NOx storage and lean NOx reduction occur simultaneously. Stored NOx is released as the temperature is elevated above the NOx storage temperature of the first component, which varies with the component's particular composition.

[0015] One embodiment of the invention system involves the use of, e.g., Pt/alumina as the first component that stores NOx at low temperatures, followed in the exhaust stream by a lean NOx catalyst that converts NOx over the higher temperatures where the Pt/alumina of the first component releases NOx. The lean NOx catalyst converts both released NOx and NOx emitted from the engine over its active temperature range. The benefit is the temperature range of the lean NOx catalyst is effectively widened to include the active temperature ranges of both components. A second embodiment of the present invention involves the use of the same two components with a modification which allows the use of the first component, e.g., Pt/alumina, for both storage and conversion of NOx. This is done by injecting HC into the exhaust to produce lean NOx reduction in the first component. (Even with the added hydrocarbons, however, the gas reacted over the first component would still be O₂-rich, usually several orders of magnitude in excess of a stoichiometric O₂ level.) In such a system, a lean NOx catalyst which follows the first component reduces released NOx. This conversion of released NOx effectively improves the level of NOx conversion for Pt/alumina. Still a third embodiment of the present invention involves a low temperature NOx storage catalyst, like Pt/alumina, with an SCR catalyst, i.e., one that uses urea or ammonia as reductant. Abatement of NOx (through absorption on the first catalyst component) will be provided over low temperatures where an SCR catalyst individually cannot operate, or where it may not be desirable to operate an SCR catalyst. As a fourth embodiment, Pt/alumina in the third embodiment could be operated similar to the second embodiment to additionally provide NOx conversion in addition to NOx storage if desired as by the addition of a source of hydrocarbons to the first component. In all of these present invention embodiment systems, using the two

components allows for the eventual reduction of NOx emitted from the engine at low temperatures. Although a single Pt-based lean NOx catalyst would be effective for converting NOx at low temperatures, however, N₂O is predominantly the product. Alternatively, when Pt/alumina is the first of the two component system of the present invention, NOx is stored at low temperatures on this first component and later reduced to the more desirable N₂ over a second component such as Cu-ZSM-5. The exhaust gas treatment system of the present invention exhibits optimal operation with diesel fuels having an absence of sulfur.

[0016] The first component, provided to absorb the nitrogen oxides present in the diesel exhaust gas at low temperatures, is particularly made of precious metal on a porous support. The porous support material is selected from the group consisting of alumina, zeolite, zirconia, titania, lanthana, and mixtures of any of them, where the alumina is preferably gamma-alumina. The support carries at least 0.1 wt. % precious metal based on the weight of the support. The precious metal is selected from the group consisting of platinum, palladium, and rhodium, or a mixture of any of them. Preferably, the precious metal is at least platinum.

[0017] The present invention NOx absorbent, contrary to those conventionally used, does not use alkali metals (e.g., potassium) or alkaline earth material (e.g., barium oxide or strontium oxide) for the absorbent. Rather, we find that the present invention support material of the first component, e.g., gamma-alumina, absorbs the NOx through the catalytic effect of the precious metal.

[0018] As described above, the support may be one or a mixture of any of the above disclosed support materials. Preferably the support is mostly gamma-alumina. By "mostly γ -alumina" in this invention is meant that the support material comprises more than about 50% by weight γ -alumina. More preferably the support is greater than 80%, and most preferably substantially all γ -alumina. In addition to the γ -alumina, however, if desired the γ -alumina support material may comprise small amounts of other materials often added into the support for stabilization of the surface area of the oxide support. This support material comprising mostly γ -alumina is desired because, in addition to being a good absorbent of NOx, it has high surface area, good adhesion and low chemical interaction with the precious metal loaded thereon.

[0019] The precious metal may be loaded onto the γ -alumina support material in any manner, several methods for providing precious metals on support materials being well known in the art. One particularly convenient method is by impregnating the support material with an aqueous or organic solvent solution comprising a soluble compound of the precious metal. Exemplary of soluble platinum (Pt) containing compounds, e.g., are compounds like chloroplatinic acid, amino-Pt-nitrates like tetramine platinum nitrate and Pt-carboxylates.

Exemplary of such solvents are water, solution of alkalis like sodium hydroxide, ethanol, toluene, isopropyl alcohol, acetone, methylethylketone, butylacetate, and dimethylformamide, water and water/methanol being particularly preferred. Any precious metal compound which is soluble in an aqueous or organic medium and whose functional group is capable of being decomposed by heat to leave only the precious metal or its oxide on the support material may be employed in this impregnation technique. Hence, such a compound is called a precious metal precursor compound. Other platinum and other precious metal precursor compounds useful in this invention in addition to those listed above will be apparent to those skilled in the art in view of the present disclosure.

[0020] According to the method of impregnation disclosed above, the precious metal precursor compound is dissolved generally by simply mixing the compound into an aqueous or organic solvent to make a precursor solution thereof. A solution of the catalyst compound is generally further diluted for impregnation of the support. The catalyst precursor solution may also comprise a mixture of compatible organic solvents and/or precursor compounds.

[0021] For useful application in an exhaust system, the washcoat will be carried on a substrate (mechanical carrier) of a high temperature stable, electrically insulating material such as cordierite, mullite, etc. A mechanical carrier is preferably comprised of a monolithic magnesium aluminium silicate structure (i.e., cordierite), although the configuration is not critical to the catalyst of this invention. It is preferred that the surface area of the monolithic structure provide 50-1000 meter square per litre structure, as measured by N₂ adsorption. Cell density should be maximized consistent with pressure drop limitations and is preferably in the range of 200-800 cells per square inch of cross-sectional area of the structure. The substrate may be in any suitable configuration, often being employed as a monolithic honeycomb structure, spun fibres, corrugated foils or layered materials. Still other materials and configurations useful in this invention and suitable in an exhaust gas system will be apparent to those skilled in the art in view of the present disclosure.

[0022] A washcoat of the support material may be applied to the substrate and then impregnated with the precious metal precursor catalyst solution. Alternately, the impregnated washcoat may be applied to the substrate. Generally, the washcoat (support material) is provided first on the substrate. Optimally, the washcoat is provided on the substrate in an amount of between about 20% and 35% by weight based on the weight of the substrate (e.g., monolith). Still other ways of providing the catalyst product will be apparent to those skilled in the art in view of the present disclosure, the method not being critical to the present invention. Preferably, a slurry of a platinum-impregnated washcoat is made into a slurry and the monolith is coated with it. When this

material is subjected to elevated temperatures the functional group is decomposed and eliminated from the catalyst precursor. It may be further subjected to calcining.

[0023] If the heating is carried out in the air, the platinum compound may exist at a platinum oxide compound which at high temperatures generally decomposes to platinum and oxygen. Thus it often exists as a mixture of platinum and its oxides.

[0024] The support material is loaded such that the amount of precious metal is at least 0.1 wt. %, more preferably, between 0.1-5.0% by weight, based on the weight of the support. Preferably when platinum is used alone without other precious metals it comprises about 0.1-5.0 wt. %, mostly preferably about 2% wt.

[0025] According to the method of the present invention, at some of the relatively low temperatures experienced during diesel engine operation, as during start up, light accels and constant speed suburban driving, the NOx in the exhaust gas will be absorbed on the first catalyst component. In particular, for a typical platinum on alumina absorbent material this will take place when the oxidizing diesel exhaust gas is 50° to 230°C. In this embodiment, when the exhaust gas temperature is raised above 230°C, generally up to about 450°C as during higher speed-load acceleration or cruise, the NOx desorbs from the absorbent and moves downstream in the exhaust gas passage. The specific temperature ranges of NOx storage and of NOx release of the first component catalyst in the present invention will depend on characteristics of its formulation as would be apparent to one skilled in the art in view of the present disclosure.

[0026] The desorbed NOx then is exposed to the second catalyst component at this elevated temperature where it is reduced, e.g. to N₂. At this elevated temperature the second component catalyst can efficiently carry out this reduction. This second catalyst component is any catalyst capable of reducing NOx, for example it may be a lean-NOx catalyst. Lean-NOx catalysts are well known in the art and are catalysts capable of reducing NOx, as well as oxidizing CO and HC, in the presence of excess oxygen. Exemplary of lean-NOx catalysts which use HC for the reduction reaction of NOx are Cu-ZSM-5, Fe-ZSM-5, Co-ZSM-5, and supported Rh. Alternately, the second component may be a SCR catalyst that reduces NOx through the use of urea or ammonia, rather than HC. Exemplary of SCR catalysts are base metal zeolites such as Cu-ZSM-5 or Ce mordenite. The invention method is not limited as to a particular catalyst for the second component as long as it performs the conversions described above. Since conventional lean-NOx catalysts reduce NOx generally by using HC in the exhaust gas, for diesel where the HC content of exhaust gas is relatively low, additional HC may be added over the lean-NOx catalyst. This may be done by injection diesel fuel, Fischer-Tropsch fuel, or other HC materials as would be apparent to those

skilled in the art in view of the present disclosure. Another way to aid reduction of the NOx when SCR catalysts are used is to introduce urea or ammonia reductant or other materials over the second component as would be apparent to those skilled in the art in view of the present disclosure.

[0027] As disclosed above, for efficient use of the system of the present invention it may be desirable to monitor the amount of nitrogen oxides absorbed by the first component at lower temperatures so as to not exceed its absorbent capacity. If the exhaust gas temperature is lower than that necessary for efficient NOx conversion by the second component, NOx can be exhausted to the atmosphere. This monitoring may be done several ways including estimating the amount of generated nitrogen oxides for engine load-time information as by an on board computer, or by measuring the amount of nitrogen oxide concentration in the exhaust gas at positions before and after the nitrogen oxide absorbent material in the passage using, e.g., a nitrogen oxide sensor. Still other ways to monitor the amount of absorbed nitrogen oxide will be apparent to those skilled in the art in view of the present disclosure. When it is determined from this monitoring that the absorbent first component trap optimally should be purged of nitrogen oxides and regenerated, the temperature of the first component is exposed to elevated temperatures which desorb the NOx. This can be done by engine calibration. There is no criticality associated with the way in which the amount of nitrogen oxides in the trap are determined nor in the way the temperature is raised to purge the absorbent. The purging is generally carried out at a selected time compatible with engine operation and for a time necessary to purge some or all of the absorbed nitrogen oxide, as selected for optimal operation of the absorbent and the engine system. It may be desirable to vary these parameters and co-ordinate them with the absorbent temperature, etc., which may aid in purging.

[0028] The stored NOx is then released from the absorbent material first component and is catalytically reduced over the catalyst of the second component catalyst. In general, the released NOx is efficiently converted to N₂ or N₂O.

[0029] Considering the alumina support, for example, our work shows that NOx is stored on the support, possibly as a nitrate. It is unstable at higher temperatures where it releases NOx. Experiments with platinum on SiO₂ showed no NOx storage to confirm the role of the alumina support as a NOx absorbent. Also, we find that NO₂, and not NO, can store and release on alumina without precious metal like platinum, but in lesser amount and over different temperature ranges than when Pt is present. The platinum appears to have two functions in the NOx storage mechanism: 1) it oxidizes NO to NO₂ which reacts with the alumina, and 2) it activates alumina in formation of an aluminium nitrate species. While this theory has been provided for an

understanding of the unexpected results of the present invention, neither the truth nor understanding thereof is necessary for a practice of the present invention.

Example 1.

Storage-Release of NOx

[0030] A cordierite honeycomb monolith, containing 33% (by weight) gamma-alumina washcoat is impregnated with chloroplatinic acid solution to deposit 2% Pt (by weight, of the washcoat). The monolith is dried at 100°C for five hours, and calcined at 500°C for four hours. This is an example of an embodiment of the first catalyst component.

[0031] The sample is tested for NOx storage and release in a flow reactor. A piece that measures 3/4" diameter x 1" long is used. Prior to testing the sample piece is reduced with a CO/N₂ feed gas and then conditioned with the test feed gas. The feed gas composition is given below.

FEED GAS COMPOSITION:

[0032]

NOx = 85 ppm
 HC = 0 ppm
 CO = 720 ppm
 H₂ = 245 ppm
 CO₂ = 5.0%
 H₂O = 4.6%
 O₂ = 14%
 N₂ = balance
 Flow rate = 3,600 cc/min

[0033] The feed gas is brought into contact with the sample at 50°C which is measured 1/2 inch before the inlet gas flow side. The flow reactor oven temperature is increased 15°C/min until the inlet sample temperature is 500°C.

[0034] The NOx storage efficiency and release as a function of inlet sample temperature is shown in Fig. 1. In this experiment, Pt/alumina shows storage over temperatures of 60-230°C. Peak storage efficiency of 80% is achieved near 160°C. NOx release is indicated by the negative values of storage efficiency that occur over temperatures of 230-450°C. The amount of NOx stored equals the amount released, within experimental error.

[0035] The stored amount of NOx is about 6 mg per in³ of catalyst monolith. This amount is below NOx storage capacity which is not reached in this experiment. However, the amount of NOx stored is on a comparable level as that which a catalyst is exposed to during the urban driving portion of the MVEuro cycle, e.g., a 2.5l diesel Transit vehicle emits ca. 2.2 g NOx which exposes a 2.5l (153 in³) catalyst to ca. 14 mg NOx per in³ catalyst. Exposure time in the experiment is 10-12

mm while NOx is storing. On the vehicle, the catalyst is exposed to exhaust for a similar time, 13-14 min, for the duration of MVEuro urban driving. It probably will not be necessary to be able to store all the engine-out NOx during the urban driving. This is because exhaust gas temperatures generally cycle through temperature ranges of both NOx storage and release.

Example 2.

Concurrent NOx Storage and NOx Reduction

[0036] The Pt/alumina monolith sample in Example 1 is tested for simultaneous occurrence of NOx storage and NOx reduction over the same temperature range. This is done by repeating the temperature ramp-up experiment in Example 1, except propylene is included in the feed gas composition. Two propylene feed gas levels are considered. They are 185 and 720 ppm HC₁, which result in 2/1 and 9/1 HC₁/NOx ratios, respectively. Both of these feed compositions with the added hydrocarbons are still oxidizing. The former represents an average level for engine-out diesel exhaust, and the latter represents a typical level when reductant is added to diesel exhaust for operating a lean NOx catalyst.

[0037] Efficiency of NOx storage plus NOx conversion as a function of inlet sample temperature is shown in Fig. 2 for 2/1 HC₁/NOx, and in Fig. 3 for 9/1 HC₁/NOx. These figures show both NOx storage and NOx conversion occurring at temperatures below 230°C. Stored NOx is released above 230°C, indicated by the negative efficiency values. The amount of NOx released is less than the amount of NOx that disappears from the feed stream for temperatures below 230°C. N₂O measurements reveal that the residual amount of NOx that disappears, i.e., the amount that is not stored, is converted, for the most part, to N₂O. For the 2/1 HC₁/NOx, NOx conversion to N₂O peaks to ca. 12-15% at 120°C, and for 9/1 HC₁/NOx, it peaks to ca. 45% at 140°C.

Example 3.

NOx Storage and Conversion of Released NOx

[0038] A two component catalyst system consisting of Pt/alumina followed by Cu zeolite (an embodiment of the invention second catalyst component) is examined in a flow reactor for NOx storage and conversion of released NOx. The Pt/alumina catalyst contacts feed gas first and is used to store NOx. This is the same catalyst sample used in Example 1. The Cu zeolite catalyst contacts feed gas after it passes through the Pt/alumina catalyst. This second position catalyst is used to convert NOx released from the Pt/alumina catalyst.

[0039] The Cu zeolite sample is a prototype lean NOx catalyst from a commercial catalyst supplier. A cordierite honeycomb monolith supports the Cu zeolite

material. Dimensions of the Cu zeolite sample are 3/4" diameter x 2" long. This sample is tested for NOx conversion with the Pt/alumina sample absent. Temperature ramp up procedure in Example 1 is performed. Feed gas composition is also the same as Example 1 except 185 ppm NOx and 1345 ppm HC₁ are used. Fig. 4 shows NOx conversion over the Cu zeolite catalyst as a function of temperature measured at the location that is inlet for the Pt/alumina catalyst but without it present. (Inlet temperature for the Cu catalyst, i.e., 1/2" in front of it, happens to be ca. 25°C higher than that in front of the Pt/alumina catalyst in this experimental apparatus.) Comparison of Fig. 4 with Fig. 1 shows NOx conversion over this Cu zeolite catalyst coincides with NOx release over Pt/alumina in the experimental set-up.

[0040] NOx conversion for the two component catalyst system, i.e., Pt/alumina followed by Cu zeolite, is shown in Fig. 5. The Pt/alumina catalyst is exposed to the same feed gas containing no HC as Example 1, so it provides NOx storage-release and no NOx conversion. Propylene is added before the Cu catalyst, i.e., after the Pt/alumina. A HC₁/NOx ratio of 15/1 is used based on the feed gas level of NO for the Pt catalyst. This high ratio is to accommodate the increase in NOx feed gas level over Cu zeolite that results from Pt/alumina releasing NOx.

[0041] Comparison of Fig. 5 with Fig. 4 shows that when the Pt/alumina storage catalyst precedes the Cu zeolite catalyst, overall net NOx conversion improves over that of the Cu zeolite alone. The improvement is from Cu zeolite converting NOx stored at the lower temperatures on Pt/alumina.

Claims

1. An exhaust gas treatment system for diesel engine exhaust, said system comprising:

a first catalyst component located in an exhaust gas passage of a diesel engine comprising a nitrogen oxide absorbent material upstream of a second catalyst component comprising a lean-NOx catalyst or a selective reduction catalyst, said nitrogen oxide absorbent material including:

- (a) porous support material selected from the group consisting of alumina, zeolite, zirconia, titania, lanthana, and mixtures of any of them; and
- (b) at least 0.1 wt. % precious metal based on the weight of the support selected from the group consisting of platinum, palladium, and rhodium, or a mixture of any of them; and

wherein said diesel exhaust gas flowing into said first component material is always oxidizing and at

low temperatures said first component material absorbs nitrogen oxides from the exhaust gas which are released from the absorbent material at higher temperatures and converted to nitrogen or nitrous oxide over the second component.

2. A system according to claim 1, wherein said precious metal comprises about 0.1-5.0 wt. %.
3. A system according to claim 1 or claim 2, wherein said precious metal comprises platinum.
4. A system according to any one of claims 1 to 3, wherein said alumina of said support comprises γ -alumina.
5. A system according to any one of the preceding claims, wherein the amount of nitrogen oxides desorbed by the first component is based on the amount of nitrogen oxides absorbed by the first component.
6. A system according to any one of the preceding claims, wherein said support material further includes stabilizers.
7. A system according to any one of the preceding claims, wherein said first component material is carried on a substrate selected from cordierite, metals or ceramic.
8. A method for treating exhaust gases generated by a diesel engine, said method comprising the steps of:

locating in an exhaust gas passage of a diesel engine a first catalyst component capable of absorbing nitrogen oxides at low temperatures and a second catalyst component downstream of said first component comprising a lean-NOx catalyst or selective reduction catalyst, said nitrogen oxide absorbent material including:

- (a) porous support material selected from the group consisting of alumina, zeolite, zirconia, titania, lanthana, and mixtures of any of them; and
- (b) at least 0.1 wt. % precious metal based on the weight of the support selected from the group consisting of platinum, palladium, and rhodium, or a mixture of any of them; and

exposing said first component material to oxidizing diesel exhaust gases, where at low temperatures nitrogen oxides are absorbed from the exhaust gas and at higher temperatures desorbed and passed to the second compo-

nent where the nitrogen oxides are reduced to nitrogen or nitrous oxide.

9. A method according to claim 8, which further comprises monitoring the absorption of the nitrogen oxide on said first component material. 5
10. A method according to claim 8 or claim 9, which further comprises injecting a source of material selected from the group consisting of hydrocarbons, ammonia or urea prior to the second catalyst component at temperatures that are at and above its light-off temperature. 10

15

20

25

30

35

40

45

50

55

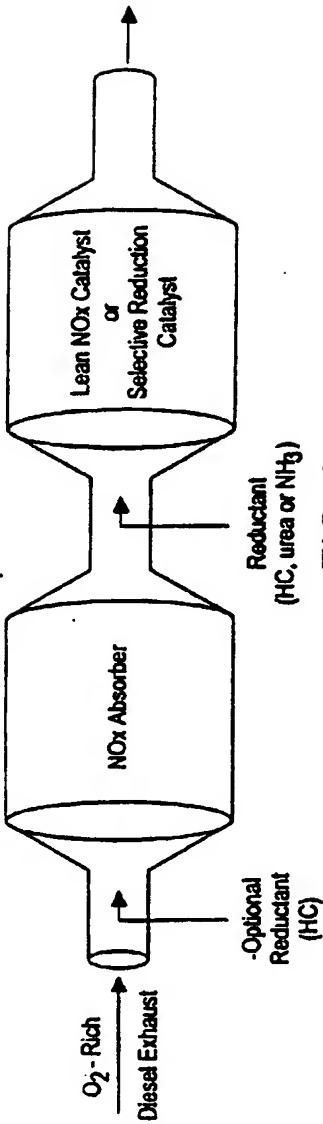


FIG. 1

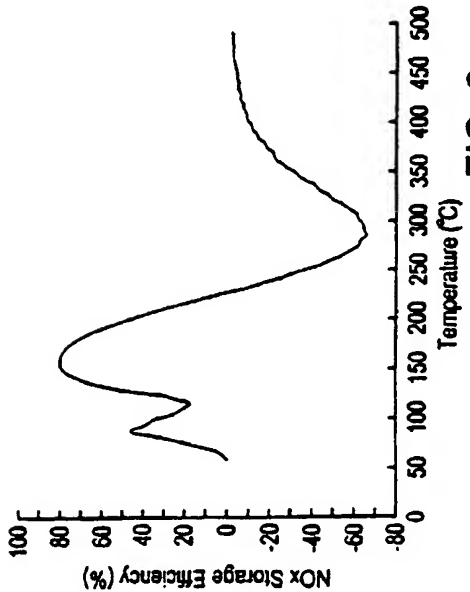


FIG. 2

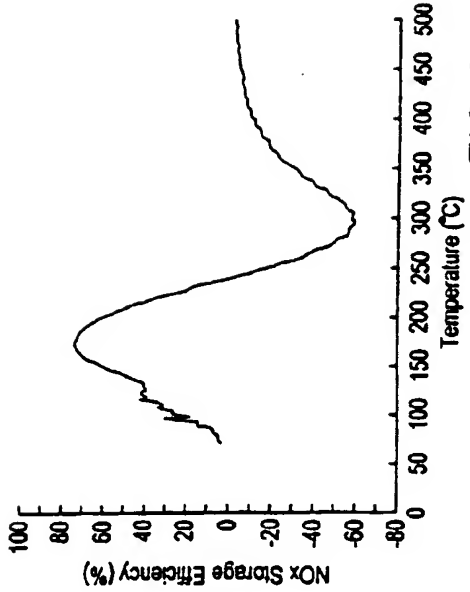
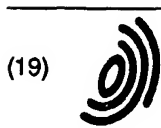


FIG. 3



(19)

Europäisches Patentamt

European Patent Office

Office européen des brevets



(11)

EP 1 027 919 A3

(12)

EUROPEAN PATENT APPLICATION

(88) Date of publication A3:
06.09.2000 Bulletin 2000/36

(51) Int. Cl.⁷: B01D 53/94, F01N 3/10,
B01J 23/40, B01J 23/42

(43) Date of publication A2:
16.08.2000 Bulletin 2000/33

(21) Application number: 00300557.6

(22) Date of filing: 26.01.2000

(84) Designated Contracting States:
AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU
MC NL PT SE
Designated Extension States:
AL LT LV MK RO SI

(72) Inventors:
• Jarvis, Mottlene
PA 19352-8933 (US)
• Adams, Karen Marie
Dearborn Heights, Michigan 48125 (US)

(30) Priority: 09.02.1999 US 248189

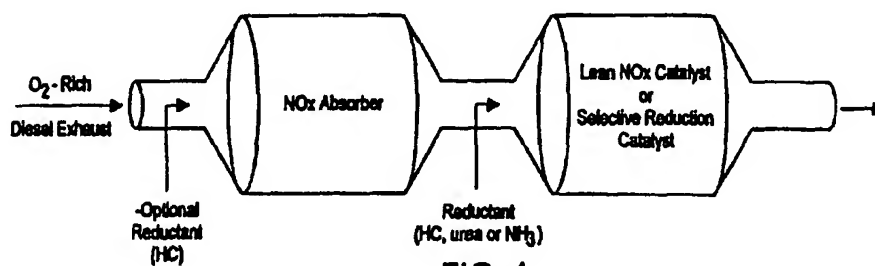
(74) Representative:
Messulam, Alec Moses et al
A. Messulam & Co. Ltd.,
43-45 High Road
Bushey Heath, Herts WD2 1EE (GB)

(71) Applicant:
Ford Global Technologies, Inc.
Dearborn, Michigan 48126 (US)

(54) Apparatus and method for treating exhaust gases from a diesel engine using nitrogen oxide absorbent

(57) The invention is a method and system for treating exhaust gases generated by a diesel engine by locating two catalyst components in the engine exhaust gas passage. The first catalyst component which is exposed to oxidizing diesel exhaust is located nearest to the engine and is a nitrogen oxide absorbent made of support material carrying precious metal. The second component is a catalyst such as a lean-NO_x catalyst or

a selective reduction catalyst which is capable of converting the exhaust gas passing over it including reducing the nitrogen oxides desorbed from the first component into nitrogen (N₂) or nitrous oxide (N₂O). Materials like hydrocarbons or ammonia or urea may be injected into the vicinity of the second catalyst component to aid in the reduction.



EP 1 027 919 A3



European Patent
Office

EUROPEAN SEARCH REPORT

Application Number
EP 00 30 0557

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.7)
X	WO 97 43031 A (ENGELHARD CORPORATION) 20 November 1997 (1997-11-20) * page 2, line 25 - page 5, line 15 * * page 13, line 14 - page 19, line 20; examples 1,2 *	1-10	B01D53/94 F01N3/10 B01J23/40 B01J23/42
X	US 5 771 686 A (PISCHINGER ET AL) 30 June 1998 (1998-06-30) * column 5, line 42 - column 6, line 13 *	1-10	
X	EP 0 886 042 A (INSTITUT FRANCAIS DU PETROLE) 23 December 1998 (1998-12-23) * column 4, line 50 - column 8, line 16 *	1-10	
X	DE 197 40 702 C (SIEMENS AG) 19 November 1998 (1998-11-19) * column 2, line 25 - column 3, line 57 *	1-10	
P,X, D	EP 0 927 571 A (FORD GLOBAL TECHNOLOGIES, INC.) 7 July 1999 (1999-07-07) * the whole document *	1-9	
P,X	EP 0 957 242 A (DAIMLERCHRYSLER AG) 17 November 1999 (1999-11-17) * column 3, paragraph 9 - column 4, paragraph 12 *	1-9	B01D F01N
P,X	DE 198 06 062 A (SIEMENS AG) 19 August 1999 (1999-08-19) * column 1, line 55 - column 3, line 1; claims 1-14 *	1-9	
A	EP 0 613 714 A (TOYOTA JIDOSHA KABUSHIKI KAISHA ET AL) 7 September 1994 (1994-09-07) * page 3, line 48 - page 6, line 37 * * page 7, line 10 - page 7, line 28 *	1-3,7,8	
		-/--	
The present search report has been drawn up for all claims			
Place of search THE HAGUE		Date of completion of the search 4 July 2000	Examiner Doolan, G
<p>CATEGORY OF CITED DOCUMENTS</p> <p>X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document</p> <p>T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document</p>			

EPO FORM 1503 03 02 (P4/C01)



European Patent
Office

EUROPEAN SEARCH REPORT

Application Number
EP 00 30 0557

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.7)
A	WO 97 02886 A (ENGELHARD CORPORATION) 30 January 1997 (1997-01-30) * page 8, line 9 - page 8, line 15 *	1,6	
A	EP 0 864 353 A (FORD GLOBAL TECHNOLOGIES, INC.) 16 September 1998 (1998-09-16) * column 4, line 11 - column 4, line 51 *	1,6	
A	DE 196 40 161 A (VOLKSWAGEN AG) 2 April 1998 (1998-04-02) * column 10, line 30 - column 11, line 62 *	8,9	
			TECHNICAL FIELDS SEARCHED (Int.Cl.7)
The present search report has been drawn up for all claims			
Place of search THE HAGUE		Date of completion of the search 4 July 2000	Examiner Doolan, G
<p>CATEGORY OF CITED DOCUMENTS</p> <p>X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document</p> <p>T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document</p>			

EPO FORM 1503 03.02 (P04001)

**ANNEX TO THE EUROPEAN SEARCH REPORT
ON EUROPEAN PATENT APPLICATION NO.**

EP 00 30 0557

This annex lists the patent family members relating to the patent documents cited in the above-mentioned European search report. The members are as contained in the European Patent Office EDP file on
The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

04-07-2000

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9743031 A	20-11-1997	AU 2430597 A	05-12-1997
		BR 9709239 A	10-08-1999
		CN 1218420 A	02-06-1999
		DE 19781766 T	08-04-1999
US 5771686 A	30-06-1998	DE 19543219 C	05-12-1996
		FR 2741385 A	23-05-1997
		GB 2307311 A, B	21-05-1997
		IT RM960768 A	08-05-1998
EP 886042 A	23-12-1998	FR 2764637 A	18-12-1998
		JP 11062567 A	05-03-1999
		US 6018943 A	01-02-2000
DE 19740702 C	19-11-1998	EP 0907010 A	07-04-1999
EP 927571 A	07-07-1999	JP 11285624 A	19-10-1999
EP 957242 A	17-11-1999	DE 19820828 A	18-11-1999
		JP 2000027634 A	25-01-2000
DE 19806062 A	19-08-1999	NONE	
EP 613714 A	07-09-1994	JP 7051544 A	28-02-1995
		JP 6262040 A	20-09-1994
		US 5911960 A	15-06-1999
WO 9702886 A	30-01-1997	US 5874057 A	23-02-1999
		AU 5422696 A	10-02-1997
		BR 9609750 A	30-03-1999
		CA 2226412 A	30-01-1997
		CN 1190909 A	19-08-1998
		EP 0850102 A	01-07-1998
		JP 11508821 T	03-08-1999
		ZA 9603196 A	04-11-1996
EP 864353 A	16-09-1998	US 5922295 A	13-07-1999
		JP 10249168 A	22-09-1998
DE 19640161 A	02-04-1998	WO 9813128 A	02-04-1998
		EP 0930930 A	28-07-1999
		US 5992142 A	30-11-1999

EPO FORM P0459

For more details about this annex : see Official Journal of the European Patent Office, No. 12/82